COMPARATIVE STUDY OF PLATELET COUNT IN DIABETIC AND NON-DIABETIC GERIATRIC POPULATION

Dr. Farheen Fatima*
Md Physiology, Upgraded Department Of Physiology, Osmania Medical College, Hyderabad, Telangana. *Corresponding Author

Dr. Ch. N. Rajkumari
Md Physiology, Prof & Hod, Upgraded Department Of Physiology, Osmania Medical College, Hyderabad, Telangana.

ABSTRACT
Geriatric population (> 60 years) is rapidly increasing in India, standing at 8.6% according to 2011 Census. Diabetes mellitus is one of the most widespread non-communicable diseases, affecting a large section of the Indian population, especially as its incidence increases with advancing age. Host of complications are associated with this disease, one of which is the effect on platelet count. This study compares platelet count between diabetic and non-diabetic elderly. It is observed that Hyperglycaemia in diabetic persons is responsible for increased Thrombopoietin production at the cellular level, which leads to raised platelet count – Reticulated Thrombocytosis – when compared to non-diabetics. Platelets, especially reticulated thrombocytes are associated with uncontrolled blood sugar levels in the body and are well known for their role in atherosclerotic Cardiovascular Disease (CVD).

INTRODUCTION:
Geriatric or elderly age group faces lot of challenges, both physiological and pathological. Aging is the progressive, universal decline first in functional reserve and then in function that occurs in organisms over time. Aging is universal but proceeds at highly variable rates, with wide heterogeneity in the emergence of the aging phenotype. Homeostatic mechanisms are slower to respond to stressors and take longer normal function. According to 2011 census, the elderly population in India is rapidly growing and stands at 8.6%, compared to 5.6% in 1961. Apart from the decline in functions which occur as part of the timescale, presence of any chronic disease adds to the morbidity in the elderly. Of the many chronic ailments, Diabetes mellitus is probably the most dreaded as it can affect multiple organ systems of our body.

Diabetes Mellitus refers to a group of common metabolic disorders that share the phenotype of hyperglycaemia. Interaction of genetic and environmental factors has led to evolution of various types of Diabetes. Based on the pathophysiology behind hyperglycaemia, Diabetes can be classified into Type 1 and Type 2.

Type 1 Diabetes is due to almost complete deficiency of insulin production whereas Type 2 Diabetes Mellitus is a heterogeneous group of disorders characterized by variable degrees of insulin resistance, impaired insulin secretion and increased glucose production. Distinct genetic and metabolic defects in insulin action and/or secretion give rise to the common phenotype of hyperglycaemia in type 2 Diabetes.

The criteria for diagnosis of Diabetes Mellitus includes a fasting blood sugar level of > 126 mg/dl (OR)
- two hours plasma glucose level of > 200 mg/dl after a glucose challenge (OR)
- Symptoms of Diabetes Mellitus (Polyuria, Polydypsia, Weight loss) + Random Blood Sugar levels of > 200 mg/dl.

Diabetes is also one of the diseases which directly affect aging systems. Conversely, both the incidence and the prevalence of diabetes mellitus increase with aging. It is a disease associated with a chronic inflammatory state in the body.

Diabetes mellitus is one of the most widespread non-communicable diseases, affecting a large section of the Indian population. 16.6% of the Indian diabetic population is concentrated solely in Hyderabad. Host of complications are associated with this disease, one of which is the effect on platelet count.

Although most standard textbooks do not point to an age-related difference in platelet count, yet some studies have observed a small, but significant decrease in platelet count of the elderly when compared to the younger age groups. However, in Diabetic individuals, this decline in platelet count is not seen, instead a rise in platelet count or thrombocytosis has been observed.

PLATELETS:
- Platelets are one of the cellular components of blood. They lack a nucleus. Also called thrombocytes. (thrombo=clot; cytes = cells).

FIGURE 1: PLATELETS – PSEUDOPODIA SEEN

Note: Although the image is red, platelets are colourless.

THROMBOPOIESIS:
Production of Platelets from the bone marrow is called Thrombopoiesis. Pluripotent Hematopoietic stem cell (HSC) converted into Megakaryocyte-Colony Forming Unit. This committed cell line develops into platelets after passing through 3 stages, which include:

1. MEGAKARYOBLAST – Earliest recognizable precursor cell, large size, has a large oval nucleus with several nucleoli.
2. PROMEGAKARYOCYTE - Formed from Megakaryoblast which undergoes ENDOREDUPPLICATION of nuclear chromatin.
3. MEGAKARYOCYTE - Promegakaryocyte matures into megakaryocyte. Single Multilobed (4-16 lobes) nucleus which contains coarsely clumped chromatin. Cell margin is irregular with many pseudopodia. Platelets are formed from detached pseudopodia containing cytoplasm and other cytoplasmic contents like granules and clotting factors etc. About 4000 platelets arise from each megakaryocyte. Formation takes about 10 days and is regulated by Colony stimulating factors and a circulating protein hormone THROMBOPOEITIN (TPO)
which increases platelet production\(^7\). TPO is produced by the liver and kidney\(^8\). Normal platelet count is between 150000 to 300000 cells per microlitre\(^9\).

**FIGURE 2: ROLE OF THROMBOPOEITIN IN PLATELET PRODUCTION**

Diabetes is a major risk factor for many systemic dysfunctions, one of which is Cardiovascular disease (CVD)\(^10\). Hyperglycaemia increases atherogenesis and atherosclerosis is one of the important causes of CVD\(^11,12\).

It has been observed that diabetic individuals have a higher platelet count when compared to non-diabetic individuals\(^13\), which is also a contributory factor in the pathogenesis of CVD\(^1\).

**AIMS AND OBJECTIVES OF THE STUDY:**
To compare the platelet count of elderly diabetics with non-diabetics.

**MATERIALS AND METHODS:**
1. **STUDY GROUP:** 30 elderly (60-70 years) Type 2 Diabetics, 13 males and 17 females, of > 5 years duration.
2. **CONTROL GROUP:** 30 age matched non-diabetics, 15 males and 15 females.

Informed Consent was taken from both groups.

**HAEMATOLOGICAL PARAMETERS ASSESSED:**
1. Platelet count – using Automated Analyzer

**INCLUSION AND EXCLUSION CRITERIA:**
**INCLUSION CRITERIA:** Subjects between 60-70 years with Type 2 Diabetes Mellitus for >5 years as study group.

Non-diabetic age matched subjects as control group.

**EXCLUSION CRITERIA:** Subjects having any other chronic diseases and/or taking drugs for the same.

**RESULTS:**
**TABLE 1: TABLE SHOWING PLATELET COUNT AND RANDOM BLOOD SUGAR LEVELS IN DIABETIC AND NON-DIABETIC INDIVIDUALS**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Mean±SD (NON-DIABETIC)</th>
<th>Mean±SD (DIABETIC)</th>
<th>p VALUES</th>
</tr>
</thead>
<tbody>
<tr>
<td>PLATELET COUNT</td>
<td>2.462 ± 0.5622</td>
<td>2.764 ± 0.5106</td>
<td>0.0365 (S)</td>
</tr>
<tr>
<td>RANDOM BLOOD SUGAR</td>
<td>119.53 ± 15.9</td>
<td>201.8 ± 41.5</td>
<td>&lt;0.0001 (ES)</td>
</tr>
</tbody>
</table>

S = Significant  
ES = Extremely Significant

Statistical analysis was done using Graphpad Instat software. Student ‘t’ test was performed.

**DISCUSSION:**
This study intended to compare the platelet count of diabetics and non-diabetic elderly individuals. RBS was done in both groups to rule out hyperglycaemia in control group. All the diabetic subjects were on oral medication only and were prescribed Metformin with or without Glimepiride by their respective Physicians. Metformin doesn’t have any effect on platelet count\(^15,16\). Side effects of Glimepiride include agranulocytosis and aplastic anaemia, though hypoglycaemia is more common. But as the cell counts of all individuals in the study group have been observed to be within normal range, it can be safely assumed that the drug didn’t interfere with the platelet count.

Both Platelet count and Random Blood sugar were observed to be significantly higher in the elderly diabetic compared to non-diabetics, although the platelet counts of both non-diabetics and diabetics were within normal range.

One of the possible explanations could be the effect of Hyperglycaemia on the increased production of IL-6\(^7\). Previous studies have shown that Hyperglycaemia leads to increased production of neutrophil derived S100 Calcium binding proteins A8/A9\(^17,18\). S100 is a multigenic family of non-ubiquitous Calcium modulated proteins of the EF – hand type family. They are implicated in intracellular and extracellular regulatory activities\(^18\). This family of proteins act as Calcium sensor proteins which participate in Calcium signal transduction by interacting with target proteins and hence modify the target’s activities\(^18\). They also mediate inflammatory responses and recruit inflammatory cells to sites of tissue damage\(^18\).

S100 proteins act as ligands for RAGE – RECEPTOR for Advanced Glycation End Products\(^18\). AGE. AGE are formed due to non-enzymatic glycosylation of intra and extracellular proteins which occur as a result of interaction of glucose with amino groups on proteins. AGEs are known to cross-link proteins such as collagen, Extracellular matrix proteins, accelerate atherosclerosis and induce endothelial dysfunction. Their serum levels correlate with degree of glycemia.\(^18\)

RAGE is seen in a variety of immune and inflammatory cells, one of them being hepatic Kupffer cells\(^19\). When S100 A8/A9 bind to the AGE receptor on the Kupffer cells, there occurs enhanced production of Interleukin-6 (IL-6). Interleukin-6, an inflammatory cytokine acts on Hepatocytes to increase production of Thrombopoetin\(^18\).

Thrombopoietin increases platelet count by acting on bone marrow progenitor cells resulting in their expansion and proliferation. It also interacts with its cognate receptor c-MPL on megakaryocytes. Ultimately reticulated thrombocytosis is
seen\(^7,12\). Reticulated platelets refers to the younger and more active platelets\(^2\). It has also been observed in previous studies that when antiplatelet therapy is prescribed to diabetic subjects with uncontrolled blood sugar levels, its usually less effective as reticulated platelets show some resistance to antiplatelet drugs\(^2,12\). Significant role of reticulated platelets have been implicated in promoting atherosclerotic plaque formation in some studies\(^2\).

Dr. A. Santakumari, Prof; Dr. R. Anitha, Prof and Dr. O. Padmini, Assoc.

### ACKNOWLEDGEMENTS:

I offer my sincere thanks to all my subjects, HOD and all the faculty of the Department of Physiology, Osmania Medical College especially the Department of Physiology, Osmania Medical College especially.

### REFERENCES:

1. WHO – Health statistics and information system; Proposed working definition of an older person in Africa for the MDS project.
2. WHO – Ageism and life course – Frequently asked questions.
5. Harrisons Principles of Internal Medicine 17th ed, Chap 33B- Diabetes Mellitus, p2275-77, 2286

### CONCLUSION:

Ageing leads to decline in functionality of the body. Added to this discomfort, there are the associated morbidities of chronic diseases, of which the most notorious one could probably be Diabetes Mellitus. Incidence of Diabetes increases with advancing age.

This study compared the platelet count of elderly diabetics with elderly non-diabetics. RBS was taken to ensure that non-diabetics had normoglycaemia as well to document the blood sugar levels in both groups. **Hyperglycaemia, was seen in the diabetic group, which lead to significant rise in platelet count in the elderly diabetic, compared to the elderly non-diabetic.** This occurred due to increased production of S100 proteins A8/A9 which are ligands for RAGE. Their interaction caused increased production of IL-6 from Hepatic Kupffer cells which caused enhanced release of Thrombopoietin from Hepatocytes. Thrombopoietin is responsible for Reticulated Thrombocytosis, which in turn is associated with complications like thrombosis, atherogenesis, CVD.

Recent studies\(^5\) have shown that if any of the steps right from production of S100 proteins till Thrombopoietin production is halted, thrombocytosis is not likely to occur. Eg: Addition of Dapagliflozin in treatment of Diabetes resulted in controlled blood sugar levels as well as absence of reticulated thrombocytosis.

Control of blood sugar levels will prevent rise of platelet count. Additionally, regular monitoring of haematological parameters like cell counts could forewarn about the development of CVD in the elderly diabetic subject.

### ACKNOWLEDGEMENTS:

I offer my sincere thanks to all my subjects, HOD and all the faculty of the Department of Physiology, Osmania Medical College especially Dr. A. Santakumari, Prof; Dr. R. Anitha, Prof and Dr. O. Padmini, Assoc.

**CONFLICT OF INTEREST:** No conflict of interest present.